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<p>(54) Title: METHODS FOR REDUCING ISCHEMIC INJURY OF THE HEART VIA THE SEQUENTIAL ADMINISTRATION OF MONOPHOSPHORYL LIPID A AND ADENOSINE RECEPTOR AGENTS</p>		
<p>(57) Abstract</p> <p>Materials and methods for reducing or preventing ischemic damage of the heart are disclosed. A preferred embodiment of the invention comprises methods of sequential administration of a plurality of cardioprotective agents to patients suffering from ischemic damage or at risk for the same.</p>		

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**METHODS FOR REDUCING ISCHEMIC INJURY OF THE HEART VIA THE
SEQUENTIAL ADMINISTRATION OF MONOPHOSPHORYL LIPID A AND
ADENOSINE RECEPTOR AGENTS**

5 Pursuant to 35 U.S.C. §202(c) it is acknowledged
that the U.S. Government has certain rights in the
invention described herein, which was made in part with
funds from the National Institutes of Health, Grant
Number HL48225.

10 **FIELD OF THE INVENTION**

The present invention relates to methods for
administering compounds to protect the heart from
ischemic injury. More specifically, the invention
provides a novel combination of agents which act
15 synergistically to potentiate their individual
cardioprotective effects and thereby render the
myocardium more resistant to ischemia.

BACKGROUND OF THE INVENTION

Several publications are referenced in this
20 application by numerals in parentheses in order to more
fully describe the state of the art to which this
invention pertains. Full citations for these references
are found at the end of the specification. The
disclosure of each of these publications is incorporated
25 by reference herein.

Ischemic preconditioning functions as an endogenous
protective mechanism which enhances the ability of
myocardium to withstand injury from prologed ischemia.
Transient ischemic events, which are sufficiently brief
30 to avoid irreversibly damaging myocardium, initiate
unidentified biochemical events which limit infarct size
and the occurrence of reperfusion arrhythmias following
prolonged myocardial ischemias (5).

A series of studies have shown that it is possible
35 to induce ischemic preconditioning with a variety of
pharmacological agents. Adenosine is released in large

amounts during myocardial ischemia and mediates potentially important protective functions in the cardiovascular system (1,4,5,7,9,14,17,18,19,25). Adenosine can precondition the heart with reduction in the size of myocardial infarction (4,5,9,14,17,18). Intracoronary administration of adenosine during reperfusion following prolonged no-flow ischemia can also limit infarct size in the intact heart (1, 19).

Previous studies have shown that adenosine A_1 and A_3 receptor agonists can precondition the heart when administered before the onset of ischemia (4, 5, 9, 14, 17, 18). Other studies have shown that adenosine A_{2a} receptor antagonists also enhance the cardioprotective effect of preconditioning (23). These agents effectively 1) reduce infarct size; and 2) improve left ventricular function when given during reperfusion (1, 19) or during both low-flow ischemia and reperfusion in isolated perfused heart (6, 21, 22).

Monophosphoryl lipid A (MLA), a relatively non-toxic derivative of endotoxin, has also been found to provide cardioprotection in a variety of animal models (43-45) when administered prior to an ischemic event. The mechanism of myocardial protection mediated by MLA has not yet been definitively elucidated. Other effective pharmacological preconditioning agents include K^+ ATP channel openers and phorbol esters.

Intensive research efforts are currently focused on the development of agents and methods for treating and preventing cardiac diseases. The present invention is directed to such methods.

SUMMARY OF THE INVENTION

Methods are disclosed which may be used to advantage for preventing ischemic damage of the heart.

The methods of the invention entail the combined administration of a first cardioprotective agent and a second cardioprotective agent. The agents act

synergistically to potentiate their individual cardioprotective effects. Agents suitable for use as a first cardioprotective agent are adenosine receptor agents, MLA or analogs or derivatives of MLA.

5 According to a preferred embodiment, the invention entails the administration of monophosphoryl lipid A (MLA), or a synthetic analog, or derivative thereof to a patient prior to a surgical treatment. Following MLA treatment, a second cardioprotective agent is
10 administered that potentiates the cardioprotective effect of MLA. The second cardioprotective agent is delivered to the patient either before, during or after surgery. Alternatively, the second cardioprotective agent may be administered continuously throughout all of these
15 periods. The protocols described above may be used for the treatment of patients at risk for ischemic damage from myocardial infarction, angina, or surgical complications.

Compounds suitable as second cardioprotective agents
20 are adenosine, agonists at the A_1 and A_3 adenosine receptors, antagonists at the A_{2a} adenosine receptor, and K_{ATP} channel openers.

In an alternative embodiment, an adenosine receptor agonist is administered as the first cardioprotective
25 agent and MLA or synthetic analogs thereof is administered as the second cardioprotective agent.

Methods of administration of the cardioprotective agents of the invention include direct perfusion of the organ during surgery and intravenous administration.
30 Additionally, these agents may be administered to patients in solid form, e.g., tablets, in amounts effective to prevent or reduce ischemic damage to the heart.

BRIEF DESCRIPTION OF THE DRAWINGS

35 Figure 1 is a graph showing the protective effect of pretreatment with MLA prior to prolonged ischemia in

cardiac myocytes.

Figure 2 is a graph showing that pretreatment with MLA enhances the cardioprotective effect of preconditioning ischemia in cardiac myocytes.

5 Figure 3A is a graph showing that pretreatment with MLA enhances the preconditioning effect mediated by adenosine in cultured myocytes (\square , vehicle; \circ , MLA 30 ng/ml; \blacksquare , MLA, 300 ng/ml). Cardiac ventricular myocytes were subjected to classical preconditioning as described
10 herein. Figures 3B and 3C are graphs showing the results obtained when myocytes were treated with CCPA (Fig. 3B) or IB-MECA (Fig. 3C) for five minutes followed by incubation in adenosine- or agonist-free media for 10 minutes prior to 90 minutes of ischemia. Data, plotted
15 as percentage of cells killed and the amount of CK released, represented the mean and standard error of four experiments. * indicates significant difference from vehicle-treated myocytes at the concentrations of adenosine agonists indicated (t test, $P < 0.001$).

20 Figure 4 is a graph showing that pretreatment with MLA enhances the preconditioning effect mediated by phorbol 12-myristate 13-acetate (PMA) (\square , vehicle; \blacksquare , MLA 300 ng/ml).

25 Figure 5 is a graph showing that pretreatment with MLA enhances the preconditioning effect mediated by pinacidil in cardiac myocytes (\square , vehicle; \circ , MLA 30 ng/ml; \blacksquare , MLA, 300 ng/ml).

30 Figure 6 is a graph showing that pretreatment with MLA enhances the cardioprotective effect of the A_1 adenosine receptor agonist, 2-chloro- N^6 -cyclopentyl-adenosine (CCPA), during prolonged ischemia (\square , vehicle; \blacksquare , MLA 300 ng/ml).

Figure 7 is a graph showing that pretreatment with MLA enhances the cardioprotective effect of the A₃ adenosine receptor agonist, N⁶-(3-iodobenzyl)adenosine-5'-N-methyluronamide (IB-MECA), during prolonged ischemia (□, vehicle; ■, MLA 300 ng/ml).

Figure 8 is a graph showing that pretreatment with increasing concentrations of MLA enhances the cardioprotective effect of pinacidil during prolonged ischemia (□, vehicle; ■, MLA 300 ng/ml).

10 DETAILED DESCRIPTION OF THE INVENTION

Preconditioning with brief ischemia before a sustained period of ischemia reduces infarct size in the perfused heart. Ventricular myocytes cultured from chick embryos retain many of the properties of the intact heart, which make them a useful model for a variety of experimental paradigms (33-39), including preconditioning (16, 23, 32). This model system has been utilized to develop novel combinations of pharmacological agents which act synergistically to treat and prevent ischemic damage of the heart.

Monophosphoryl lipid A (MLA), a derivative of the gram-negative bacterial cell wall component lipopolysaccharide A (i.e. endotoxin), induces a glibenclamide-sensitive cardioprotective effect in isolated, intact heart models of ischemia-reperfusion. The data reveal that treatment with MLA has a prolonged protective effect (31). The present invention is based on the hypothesis that MLA or MLA analogs effect cardioprotective action by enhancing the sensitivity of the cardiac myocytes to preconditioning stimuli and to cardioprotective agents. These cardioprotective agents include agonists at the A₁ and A₃ adenosine receptors, antagonists at the A_{2a} adenosine receptor, K_{ATP} channel openers and phorbol esters. MLA may also enhance the

protective effects of these agents during prolonged, infarct-producing ischemia. These two novel cardioprotective mechanisms of MLA were investigated and the results are provided hereinbelow.

5 The present invention demonstrates that pretreatment of myocytes with MLA alone prior to preconditioning ischemia can enhance the cardioprotective effect of the preconditioning, thus further decreasing myocyte killing in response to an ischemic event.
10 Furthermore, a synergistic cardioprotective effect is achieved by treatment with MLA in conjunction with adenosine, agonists of the A₁ and A₃ adenosine receptors, antagonists of the A_{2a} adenosine receptor or a K_{ATP} channel opening compound.

15 The compounds of the present invention tend to produce certain unwanted side effects when administered individually at standard dosage. Because these compounds have been discovered to act synergistically when used in combination, subthreshold concentrations may be employed
20 in practicing the methods of the present invention. These subthreshold concentrations should induce fewer side effects and accordingly constitute an improvement in currently available methods for reducing ischemic damage to the heart.

25 The following definitions are provided to facilitate understanding of the present invention.

Preconditioning ischemia - A brief ischemia which does not cause any cardiac damage, but protects the heart against damage during a subsequent prolonged ischemia.

30 The effect of preconditioning ischemia is mediated by adenosine, which is released during the ischemia. Preconditioning may be induced by brief exposure to anoxic conditions for example.

Adenosine receptors - A₁, A₃ and A_{2a} receptors are present
35 on the myocardium (cardiac muscle cells). While activation of the A₁ and A₃ receptors is cardioprotective, activation of the A_{2a} receptors is deleterious and causes

damage to the cardiac muscle cells.

Stable angina - Condition observed in certain cardiac patients having a chronic risk for myocardial ischemia because of the chronic potential for an imbalance between the supply of oxygenated blood and the demand for it. Typically, such imbalance occurs during certain stresses, such as exercise, emotional stress or stress associated with a surgical procedure.

Unstable angina - Condition observed in cardiac patients having frequent imbalance between the supply of and the demand for oxygenated blood.

Post-myocardial infarction angina - Condition observed in patients who have recurrent ischemia following a heart attack.

Preconditioning stimuli - Any drug, agent or treatment which induces preconditioning, such as brief hypoxia, adenosine, pinacidil, phorbol ester, A_1 or A_3 adenosine receptor agonists or A_{2a} receptor antagonists.

Myocardial responsiveness - The myocardium can be treated so as to enhance the effectiveness and protective effects of preconditioning. This enhancement leads to a reduction in ischemic damage.

The following methods facilitate the practice of the present invention.

25 Preparation of Cultured Ventricular Cells

Ventricular cells were cultured from chick embryos 14 days in ovo (Spafas Inc., Storrs, CT) as previously described (16, 23). Cells were cultivated in a humidified 5% CO_2 -95% air mixture at 37 C. All experiments were performed on day 3 in culture, at which time cells exhibited rhythmic spontaneous contraction. The medium was changed to a HEPES-buffered medium containing (mM) 139 NaCl, 4.7 KCl, 0.5 $MgCl_2$, 0.9 $CaCl_2$, 5 HEPES (N-2-hydroxyethylpiperazine-N'-2-ethanesulfonic acid) and 2% fetal bovine serum, pH 7.4, 37 C before

exposing the myocytes to the various conditions at 37 C. Control myocytes were maintained in HEPES-buffered medium under room air. Hypoxia and glucose deprivation were used to induce and simulate ischemia in the cultured
5 cells. Hypoxia was produced by placing the cells in a hypoxic incubator (NuAire) where O₂ was replaced by N₂ as previously described (16,23).

Determination of Cell Injury

The extent of hypoxia-induced injury to the
10 ventricular cell was quantitatively determined by the percentages of cells killed. To quantitate the % of cells killed, cells were exposed to a trypsin-EDTA Hanks' balanced salt solution for 10 minutes for detachment
15 after the various treatment, followed by centrifugation (300 x g for 10 minutes) and resuspension in culture media for counting in a hemocytometer. Only live cells sedimented and the cells counted represented those that survived (46). None of the sedimented cells subsequently
20 counted included trypan blue. In cells not exposed to hypoxia, trypsin-EDTA treatment followed by re-exposure to Ca²⁺-containing culture media did not cause the appearance of trypan blue-stained cells or any
25 significant increase in proteins or creatine kinase (CK) in the trypsin-EDTA media following the 300 x g, 10 minute sedimentation of the cells. There was no protein
30 in the culture media following a second 300 x g centrifugation of resuspended cells previously treated with trypsin-EDTA. Thus, trypsin treatment, re-exposure to Ca²⁺-containing media or 300 x g sedimentation did not cause any significant damage to the control,
normoxia-exposed cells.

The percentage of cells killed was calculated as the number of cells obtained from the control group
(representing cells not subjected to any hypoxia or drug
35 treatment) minus the number of cells from the treatment group divided by number of cells in control group

multiplied by 100%.

Chemical reagents

2-chloro-N⁶-cyclopentyl-adenosine (CCPA) and pinacidil were purchased from Research Biochemicals International (Natick, MA). Phorbol 12-myristate 13-acetate (PMA) was
5 purchased from Sigma. 8-(3-chlorostyryl)caffeine (CSC), N⁶-(3-iodobenzyl)adenosine-5'-N-methyluronamide (IB-MECA) and 2-chloro-N⁶-(3-iodobenzyl)adenosine-5'-N-methyluronamide (Cl-IB-MECA) were synthesized as
10 previously described (40-42).

The following examples are provided to illustrate certain embodiments of the invention. They are not intended to limit the invention in any way.

EXAMPLE I

15 A cultured chick ventricular myocyte model was developed to investigate the roles of a variety of compounds known to influence preconditioning. In accordance with the present invention, it has been discovered that certain agents, when applied in
20 combination, act synergistically to reduce ischemic damage to the heart.

Pretreatment of ventricular myocytes with MLA was assessed to determine whether it enhances the cardioprotective effect of preconditioning ischemia.
25 Hypoxia (O₂ <1%) and glucose deprivation were employed to induce and simulate ischemia in the myocyte model of preconditioning (16, 23, 32).

Cardiac ventricular myocytes were cultured from chick embryos 14 days in ovo and myocyte injury was
30 induced as described hereinbelow. MLA was added to the media at the indicated concentrations for a 4 hour period 24 hours prior to 90 minutes of simulated ischemia. The extent of myocyte injury, determined at the end of the 90 minute simulated ischemia, was assessed by the percentage

of cells killed. The data represent the mean of 4 experiments. At 30 and 300 ng/ml MLA significantly lowered the percentage of cell death in response to prolonged ischemia. (One way ANOVA analysis and t test, $p < 0.01$).

Exposure of cultured cardiac myocytes to 90 minutes of prolonged hypoxia causes cell death. Figure 1 illustrates that prior treatment with MLA significantly reduces cell death relative to the untreated controls. These data are in agreement with the cardioprotective effect of MLA treatment seen in intact rat, rabbit, and canine hearts (43-45).

Figure 2 shows that the extent of cardioprotection induced by a 5 minute period of preconditioning ischemia (induced by hypoxia and the absence of glucose, also termed simulated ischemia) is enhanced by pretreatment with MLA. As in Figure 1, cultured myocytes were pretreated with the indicated concentrations of MLA for 4 hours. Twenty four hours later the cells were subjected to preconditioning ischemia, consisting of 5 minutes of simulated ischemia. After a 10 minute recovery period in normal air, the cells were subjected to a prolonged period (90 minutes) of simulated ischemia, and the percentage of cells killed was assessed. When myocytes are pretreated with MLA 24 hours prior to preconditioning ischemia treatment, there is a dose-dependent reduction in the percentage of cells killed during a prolonged ischemic event. Thus, pretreatment with MLA in combination with classic preconditioning ischemia provides a synergistic cardioprotective effect.

EXAMPLE II

Another objective of this study was the elucidation of the mechanism underlying the observed enhanced responsiveness of cardiac myocytes to preconditioning stimuli following pretreatment with MLA. The cardioprotective effects of preconditioning ischemia are mediated by adenosine receptors and the K_{ATP} channel. The

following experiments were performed to assess the most efficacious combination of agents for the prevention of ischemic damage to the heart.

As shown in Figure 3A, prior treatment with MLA enhanced the ability of adenosine to elicit the preconditioning response. In this experiment, myocytes were treated with either the vehicle (0.04% propylene glycol, 0.01% ethyl alcohol) or MLA 24 hours prior to a 5 minute exposure to 10 μ M adenosine. Adenosine was removed by replacement of the culture medium with adenosine-free medium, and the cells were exposed to normal room air O₂ for 10 minutes, followed by 90 minutes of simulated ischemia treatment. The sequential treatment of myocytes with MLA followed by adenosine significantly decreased the percentage of myocytes killed by prolonged ischemia, as compared to pretreatment with the vehicle followed by exposure to adenosine. Similarly, Figure 3B and 3C show that prior treatment of myocytes with MLA followed by adenosine agonist, CCPA or by A3 agonist, IB-MECA, as a classical preconditioning stimuli, also significantly reduced the percentage of myocytes killed by prolonged ischemia as compared to pretreatment with the vehicle followed by exposure to CCPA or IB-MECA alone, respectively.

An enhanced adenosine-mediated classical preconditioning response in myocytes pre-exposed to MLA suggests that MLA acts to increase the responsiveness of myocytes to the preconditioning effect of adenosine. Since the adenosine receptor mediates the preconditioning response to adenosine, such results may further suggest that MLA activates or pre-activates the adenosine receptor/G-protein/PKC/K_{ATP} channel pathway. Consistent with this hypothesis, the ability of the phorbol ester phorbol 12-myristate 13-acetate (PMA) to mediate a preconditioning response was also enhanced by pretreatment with MLA. See Figure 4. This enhancement by 300 ng/ml MLA pretreatment was significant over a

range of PMA concentrations from 0.01 μ M to 1 μ M. The preconditioning effects of CCPA or IB-MECA were also enhanced by pretreatment of myocytes with MLA. This enhancement was not observed in myocytes pretreated with vehicle alone (data not shown).

MLA pretreatment also enhanced the preconditioning response mediated by pinacidil. See Figure 5. The data are consistent with the notion that MLA can pre-activate the adenosine receptor, G protein, PKC, the K_{ATP} channel, or all four molecules. On the other hand, as the K_{ATP} channel is the most distal effector of the four, MLA may pre-activate or activate the K_{ATP} channel directly to render the cardiac myocyte more responsive to the preconditioning effect of the K_{ATP} channel opener pinacidil.

EXAMPLE III

The experiments described in the previous examples suggest that MLA acts to enhance the preconditioning effects of various pharmacological agents such as adenosine and pinacidil.

The synergistic effects obtained using a combination of MLA and other preconditioning stimuli prompted studies assessing the activity of agonists and antagonists of adenosine receptors following pretreatment with MLA.

The experimental protocol utilized in these studies entailed pretreatment of myocytes with MLA for a 4 hour period 24 hours prior to exposure of the cardiac myocytes to 90 minutes of simulated ischemia. During this 90 minute simulated ischemia period, several cardioprotective agents were administered individually to cells and their protective effects were assessed.

Figure 6 illustrates that prior treatment of myocytes with MLA enhanced adenosine A_1 agonist mediated protection during the 90 minute exposure to ischemia. In MLA pretreated myocytes, the presence of CCPA during the 90 minute ischemia resulted in a greater reduction in the

number of cells killed as compared to the number of cells killed following pretreatment with vehicle alone. Again the synergistic effects mediated by the two cardioprotective agents are illustrated by the data.

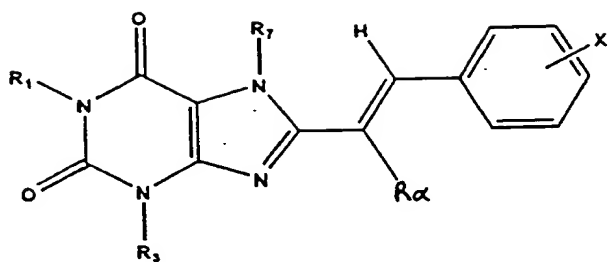
5 Similar results were obtained when cells were pretreated with MLA then treated with IB-MECA, an A₃ adenosine receptor agonist during the 90 minute ischemia as described above for Figure 6. See Figure 7. Again the results showed a synergistic cardioprotective effect
10 mediated by the combination of the two agents.

Figure 8 shows the results obtained by pretreatment with MLA followed 24 hours later, by exposure to pinacidil, a K_{ATP} channel opener, during the 90 minute ischemia. In MLA-pretreated myocytes, pinacidil caused a
15 greater reduction in the number of cells killed as compared to vehicle pretreatment.

Additional experiments were performed to assess the differences in cardioprotection observed when MLA was administered as the second cardioprotective agent and an
20 adenosine agonist, KATP channel opener or protein kinase C activator was administered as the first cardioprotective agent. The experiments were performed as follows: Cardiac myocytes were exposed to the first cardioprotective agent for 5 minutes and the agent was
25 removed. Cells were then exposed to a period of simulated ischemia in the presence or absence of 300 ng/ml of MLA. The percentage of cells killed by simulated ischemia alone was $27 \pm 2\%$ (n=9) in the absence of any treatment with any cardioprotective agent. When
30 MLA was present during the simulated ischemia, the percentage of cells killed was reduced from $27 \pm 2\%$ to $14 \pm 1.2\%$. When cells were exposed to 5 minutes of $10 \mu\text{M}$ pinacidil, a KATP channel opener followed by exposure to simulated ischemia (in the absence of MLA), the
35 percentage of cells killed was 12.3 ± 1 . When the pinacidil-treated cells were exposed to simulated ischemia in the presence of MLA, the percentage of cells

killed was reduced from $12.3 \pm 1\%$ to $6.3 \pm 1.2\%$ ($n=3$).
When cells were exposed to 5 minutes of $1 \mu\text{M}$ PMA, a PKC
activator, followed by exposure to simulated ischemia
(in the absence of MLA), the percentage of cells killed
5 was 14 ± 1 . When PMA-treated cells were then exposed to
simulated ischemia in the presence of MLA, the percentage
of cells killed was reduced to 10.5 ± 1 ($n=3$). When cells
were exposed to 5 minutes of $10 \mu\text{M}$ adenosine, followed by
exposure to simulated ischemia (in the absence of MLA),
10 the percentage of cells killed was $12 \pm 1.4\%$. When
adenosine treated cells were then exposed to simulated
ischemia, in the presence of MLA, the percentage of cells
killed was reduced to $7.4 \pm 1.2\%$. These results indicate
that the cardioprotective agents of the invention are
15 effective in reducing ischemic damage of the heart
regardless of the order in which they are administered to
cells.

The data presented in the Figures illustrate the
synergistic effects mediated by the combination of MLA,
20 adenosine receptor agonists and K_{ATP} channel openers in
mediating cardioprotection. Accordingly, other agents
known to act on the adenosine receptors are contemplated
for use in the present invention. Earlier studies have
revealed that antagonists at the A_{2a} receptor mediate
25 cardioprotective effects. Thus, the combined use of MLA
or analogs thereof followed by treatment with an A_{2a}
antagonist should also prove to be efficacious in
reducing ischemic injury of the heart. Candidate
compounds of the following formula that bind and inhibit
30 adenosine A_{2a} receptor activation are set forth below in
Table I.



R₁, R₃ = methyl, ethyl, propyl, allyl

R₇ = H, alkyl (C1-C8)

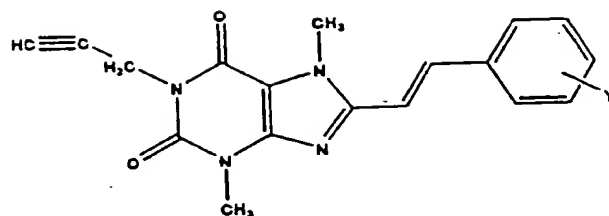
R_α = H

TABLE I
Affinities of 8 Styrylxanthine Derivatives in
Radioligand Binding Assays at Rat Brain A₁ and A₂-Receptors (47)

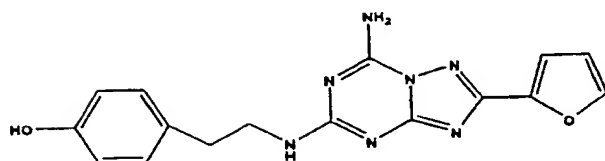
Radiolabeled Compounds					
compd	R ₁ , R ₂	R ₃	X	A ₁ /A ₂ ratio	
5	15b	Me	Me	H	41
	17b	Me	Me	2-MeO	18
	19b	Me	Me	3-MeO	64
	20b	Me	Me	3-CF ₃	25
	21b	Me	Me	3-NO ₂	11
10	22b	Me	Me	3-NH ₂	30
	23	Me	Me	3-(AcNH)	240
	24	Me	Me	3-(HOOC(CH ₂) ₂ CONH)	250
	25	Me	Me	3-(t-BOC-NH)	30
	26	Me	Me	3-[t-BOC] ₂ N]	15
15	27b	Me	Me	3-F	190
	28	Me	Me	3-Cl	520
	29b	Me	Me	4-MeO	44
	32b	Me	Me	3,4-(MeO) ₂	70
	33a	Me	H	3,5-(MeO) ₂	25
20	33b	Me	Me	3,5-(MeO) ₂	>200
	34b	Me	Me	3,5-F ₂	230
	35	Me	Me	3,5-(MeO) ₂ -4-OH	19
	36	Me	Me	4-AcO-3,5-(MeO) ₂	93
	37	Me	Me	4-(4-PhCH ₂ O)-3,5-(MeO) ₂	30
25	38	Me	Me	4-(4-NH ₂ -BuO)-3,5 (MeO) ₂	36
	39	Me	Me	4-[4-(tBOC-NH) BuO]-3,5-(MeO) ₂	42
	40	Me	Me	4-(4-NH ₂ -trans-CH ₂ CH=CHCH ₂ O-3,5-(MeO) ₂	28
	41	Me	Me	4-(4-AcNH-trans-CH ₂ CH=CHCH ₂ O)-3,5-(MeO) ₂	>50
	42	Me	Me	4-(4-t-BOC-NH-trans-CH ₂ CH=CHCH ₂ O-3,5-(MeO) ₂	>40
30	43b	Me	Me	2,3,4-(MeO) ₃	34
	44b	Me	Me	3,4,5-(MeO) ₃	70 [>5600]
	45b	Et	Me	3,4,5-(MeO) ₃	34
	46	allyl	Me	3,4,5-(MeO) ₃	13 [>6700]
	51b	Pr	Me	3-Cl	14
35	52b	Pr	Me	3,4-(MeO) ₂	19 [190]
	53b	Pr	Me	3,5-(MeO) ₂	110

Additional compounds contemplated for use as A_{2a}

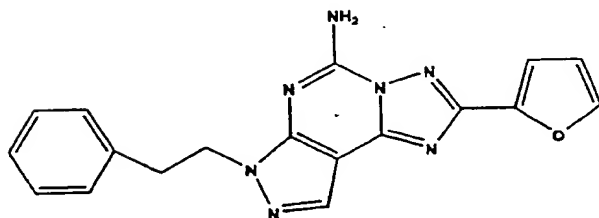
antagonists include:



Y = m-Br or p-SO₃H (DMPX derivative)



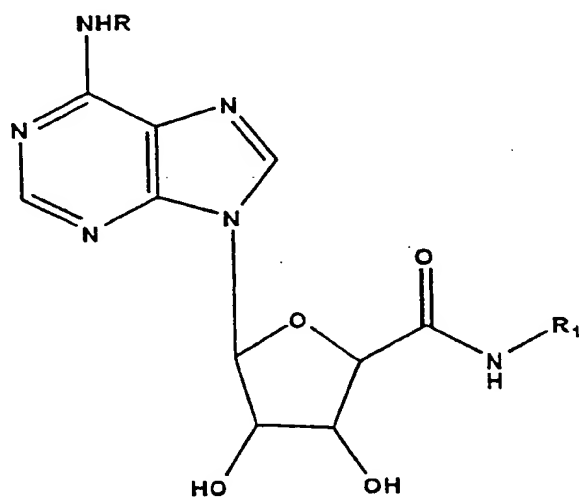
5 (ZM241385)

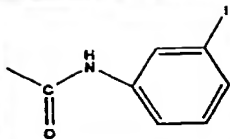
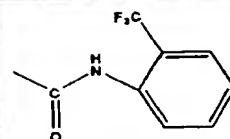
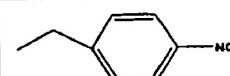
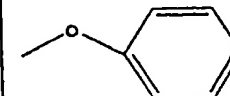
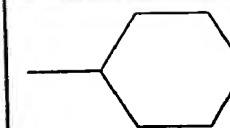


(SCH58261)

As mentioned previously, agonists at the A₁ and A₂ adenosine receptors are also contemplated for use in the methods of the present invention. Suitable agonists of the following formula are set forth below in Table II.

5



COMPOUND (Reference)	R ₁	R	A ₁ (nm)	A _{2a} (nm)	A ₃ (nm)
Compound 4q (Reference 27)	Ethyl		16	3940	30
Compound 4d (Reference 27)	Ethyl		384	>10,000	54
Compound 37 (Reference 8)	Ethyl		49	574	9.0
Compound 11 (Reference 28)	Methyl		2060	66,300	1340
N ⁶ -cyclohexyl NECA (Reference 29)	Ethyl		0.43	170	16

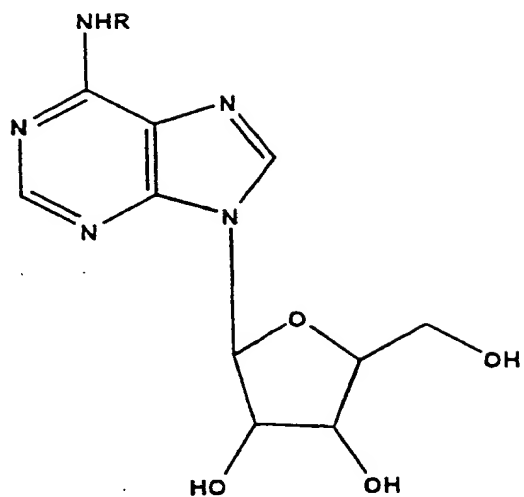
4q = N⁶-((3-iodophenyl)carbamoyl)adenosine -5'uronamide

4d = N⁶-((2-trifluoromethyl)carbamoyl)adenosine-5'uronamide

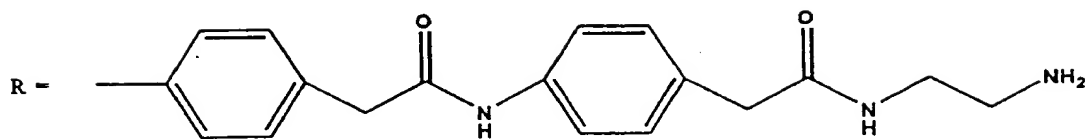
compound 37 = N⁶-(4-Nitrobenzyl)adenosine-5'-N-ethyluronamide

compound 11 = 6-(O-Phenylhydroxylamino)purine-9-beta-ribofuranoside-5'-N-methyluronamide

N⁶-cyclohexyl NECA = N⁶-cyclohexyl 5'-N-ethylcarboxamidoadenosine



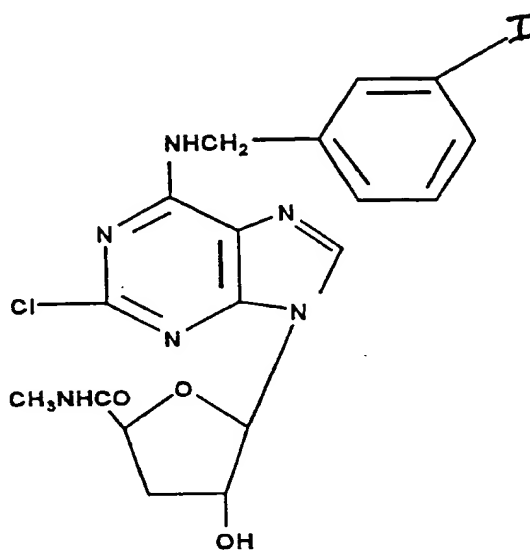
- O, N, R



COMPOUND	R	A ₁ (nm)	A ₂ (nm)	A ₃ (nm)
Compound 8'	(See immediately above.)	0.85	210	4

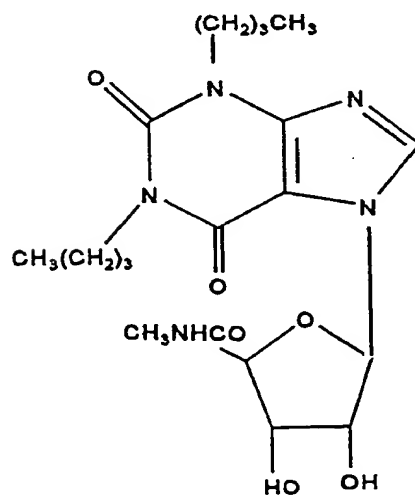
5 *N⁶-[4-[[[4-[2-aminoethyl)amino]carbonyl]methyl]-
anilino]carbonyl]methyl]phenyl]adenosine

Other effective A3 agonists suitable for use in practicing the methods of the present invention are set forth below:

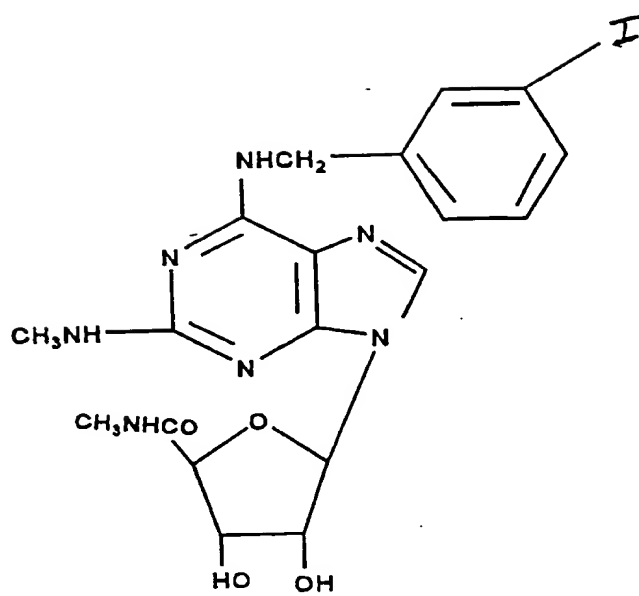


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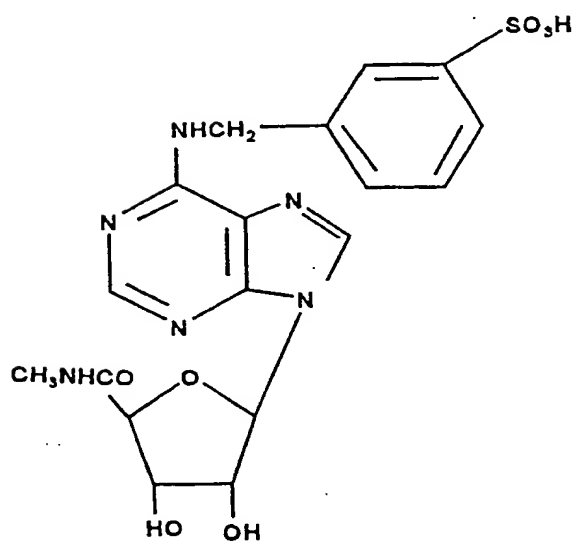
MRS 584



MRS 479 (48)



MRS 537 (13)



MRS 1340 (8)

EXAMPLE IV

The synergistic cardioprotective effect of MLA used in combination with A_3/A_1 agonists and a K_{ATP} channel opener has been demonstrated herein. Combinations of these agents may be used therapeutically in patients who suffer from ischemic damage due to stable angina, unstable angina or post-myocardial infarction angina.

Several administration modalities may be utilized to treat patients with the agents of the invention. These modalities are influenced by bioavailability factors. For example, if the compound is metabolized in the liver or excreted in the bile, some of the active compound absorbed from the gastrointestinal tract will be inactivated by the liver before it can reach the general circulation and for distribution to the site of action. It is not believed that adenosine A_1/A_3 receptor agonists, adenosine A_{2a} receptor antagonists or K_{ATP} will be subject to this first pass loss. Additionally, because the agonists of the invention are polar and water soluble, it is expected that they will have a small volume of distribution, and thus be readily eliminated by the kidney. Moreover, binding of these agents to plasma proteins may limit their free concentrations in tissues and at their locus of action since it is only the unbound drug which equilibrates across membrane receptor sites.

Another factor affecting bioavailability is the distribution of the agonists to tissues. Given the relatively small size of these compounds and their water solubility, it is anticipated that the compounds will have a relatively fast second phase of drug distribution. This distribution is determined by both the blood flow to the particular tissue of the organ such as the heart, as well as the rate at which the compounds diffuse into the interstitial compartment from the general circulation through the highly permeable capillary endothelium.

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MLA or synthetic analogs thereof are lipophilic and will be administered intravenously in a 10% ethanol, 40% propylene glycol water solution. Alternatively MLA or its analogs can be given orally in capsules or tablets. 5 MLA or its analogs would be administered in doses ranging from 1-20 $\mu\text{g/kg}$ body weight. Patients may be perfused with the adenosine receptor A_1 or A_3 agonists, adenosine A_{2a} receptor antagonists or K_{ATP} channel openers of the invention by dissolving them in normal saline solution or 10 using emulsifying agents or cosolvents followed by intravenous administration every four to six hours. Effective doses usually range from 100 to 300 nM. For example, considering a 15 liter volume of distribution for a 70 kg patient, a loading dose ranging from 0.5 to 15 1.5 mg is preferably used. Depending on the half-life of the agonists in the body, several doses, e.g., 1.5-4.5 mg may be administered per day.

Alternatively, a time-release or slow-release preparation may be utilized which allows for periodic or 20 constant release of the agonists, antagonists or channel openers over a given time period. This method would allow for a single dose of these agents in a given day. Methods for preparing such capsules are well known to those of skill in the art of drug delivery.

25 While the combined use of MLA with specific A_3 and A_1 agonists or K_{ATP} channel openers has been exemplified herein, other compounds have been identified which have high affinity for both the A_3 and A_1 receptor, also other compounds are known to those skilled in the art that 30 function to open the K_{ATP} channel. Additionally, analogs or derivatives of MLA are known. Any of these compounds or agents therefore may also be used in the practice of the instant invention.

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While certain preferred embodiments of the present
20 invention have been described and specifically
exemplified above, it is not intended that the invention
be limited to such embodiments. Various modifications
may be made to the invention without departing from the
scope and spirit thereof as set forth in the following
25 claims.

What is claimed is:

1. A method for preventing or reducing ischemic heart damage, in a patient in need of such treatment, comprising administering to said patient a first
5 cardioprotective agent followed by administration of a second cardioprotective agent, said first and second cardioprotective agents acting synergistically to protect the heart from ischemic damage.
2. A method as claimed in claim 1, wherein said
10 first cardioprotective agent is selected from the group consisting of monophosphoryl lipid A, a synthetic analog of monophosphoryl lipid A, adenosine A1 agonists or adenosine A3 agonists.
3. A method as claimed in claim 1, wherein said
15 second cardioprotective agent is selected from the group consisting of adenosine A1 receptor agonists, adenosine A3 receptor agonists, agonists listed in Table II of the present specification, MRS 584, MRS 479, MRS 537, MRS 1340, adenosine A2a receptor antagonists, antagonists
20 listed in Table I of the present specification, PKC activators and K_{ATP} channel openers.
4. A method as claimed in claim 1, wherein said
25 first cardioprotective agent is administered to said patient by an administration means selected from the group consisting of intravenous administration, direct cardiac perfusion, or oral administration.
5. A method as claimed in claim 1, wherein said
30 second cardioprotective agent is administered to said patient by an administration means selected from the group consisting of intravenous administration, direct cardiac perfusion, or oral administration.

6. A method as claimed in claim 1, wherein said patient is in need of said treatment due to the presence of angina selected from the group consisting of unstable
5 angina or post-myocardial infarction angina.

7. A method as claimed in claim 1, wherein said cardioprotective agents are administered to said patient prior to a surgical procedure which may cause cardiac ischemic damage.

10 8. A method as claimed in claim 1, wherein said cardioprotective agents are administered to said patient during a surgical procedure which may cause cardiac ischemic damage.

15 9. A method as claimed in claim 1, wherein said cardioprotective agents are administered to said patient following a surgical procedure which may cause cardiac ischemic damage.

20 10. A method for enhancing myocardial responsiveness to preconditioning stimuli in a patient in need of such treatment comprising administering to said patient a first cardioprotective agent followed by administration of a second cardioprotective agent, said first and second cardioprotective agents acting synergistically to enhance myocardial responsiveness.

25 11. A method as claimed in claim 10, wherein said first cardioprotective agent is selected from the group consisting of monophosphoryl lipid A, a synthetic analog of monophosphoryl lipid A, adenosine A1 agonists or adenosine A3 agonists.

30 12. A method as claimed in claim 10, wherein said second cardioprotective agent is selected from the group

consisting of adenosine A1 receptor agonists, adenosine A3 receptor agonists, adenosine A2a receptor antagonists, antagonists listed in Table I of the present specification, PKC activators and K_{ATP} channel openers.

5 13. A method as claimed in claim 10, wherein said first cardioprotective agent is administered to said patient by an administration means selected from the group consisting of intravenous administration, direct cardiac perfusion, or oral administration.

10 14. A method as claimed in claim 10, wherein said second cardioprotective agent is administered to said patient by an administration means selected from the group consisting of intravenous administration, direct cardiac perfusion, or oral administration.

15 15. A method as claimed in claim 10, wherein said patient is in need of said treatment due to the presence of angina selected from the group consisting of unstable angina or post-myocardial infarction angina.

20 16. A method as claimed in claim 10, wherein said cardioprotective agents are administered to said patient prior to a surgical procedure which may cause cardiac ischemic damage.

25 17. A method for reducing or preventing ischemic heart damage in a patient in need of such treatment, comprising the administration of first and second cardioprotective agents, said first cardioprotective agent being selected from the group consisting of A1 adenosine agonists or A3 adenosine agonists, K_{ATP} channel
30 openers, and PKC activators followed by the administration of a second cardioprotective agent selected from the group consisting of monophosphoryl lipid A or synthetic analogs of monophosphoryl lipid A.

18. A method as claimed in claim 17, wherein said cardioprotective agents are administered to said patient prior to a surgical procedure which may cause cardiac ischemic damage.

5 19. A method as claimed in claim 17, wherein said cardioprotective agents are administered to said patient during a surgical procedure which may cause cardiac ischemic damage.

10 20. A method as claimed in claim 17, wherein said cardioprotective agents are administered to said patient following a surgical procedure which may cause cardiac ischemic damage.

15 21. A method as claimed in claim 17, wherein said first cardioprotective agent is administered to said patient by an administration means selected from the group consisting of intravenous administration, direct cardiac perfusion, or oral administration.

20 22. A method as claimed in claim 17, wherein said second cardioprotective agent is administered to said patient by an administration means selected from the group consisting of intravenous administration, direct cardiac perfusion, or oral administration.

25 23. A method as claimed in claim 17, wherein said patient is in need of said treatment due to the presence of angina selected from the group consisting of unstable angina or post-myocardial infarction angina.

24. A method as claimed in claim 17, wherein said K_{ATP} channel opener is nicorandil.

30 25. A method as claimed in claim 17, wherein said first cardioprotective agent is selected from the group

consisting of agonists listed in Table II of the present specification, MRS 584, MRS 479, MRS 537, and MRS 1340.

26. A method for enhancing myocardial responsiveness to preconditioning stimuli in a patient in need of such treatment comprising administering to said patient a first cardioprotective agent followed by administration of a second cardioprotective agent, said first cardioprotective agent being selected from the group consisting of A1 adenosine agonists or A3 adenosine agonists, K_{ATP} channel openers, and PKC activators followed by the administration of a second cardioprotective agent selected from the group consisting of A2a antagonists, monophosphoryl lipid A or synthetic analogs of monophosphoryl lipid A, said first and second cardioprotective agents acting synergistically to enhance myocardial responsiveness.

27. A method as claimed in claim 26, wherein said first cardioprotective agent is administered to said patient by an administration means selected from the group consisting of intravenous administration, direct cardiac perfusion, or oral administration.

28. A method as claimed in claim 26, wherein said second cardioprotective agent is administered to said patient by an administration means selected from the group consisting of intravenous administration, direct cardiac perfusion, or oral administration.

29. A method as claimed in claim 26, wherein said patient is in need of said treatment due to the presence of angina selected from the group consisting of unstable angina or post-myocardial infarction angina.

30. A method as claimed in claim 26, wherein said cardioprotective agents are administered to said

patient prior to a surgical procedure which may cause cardiac ischemic damage.

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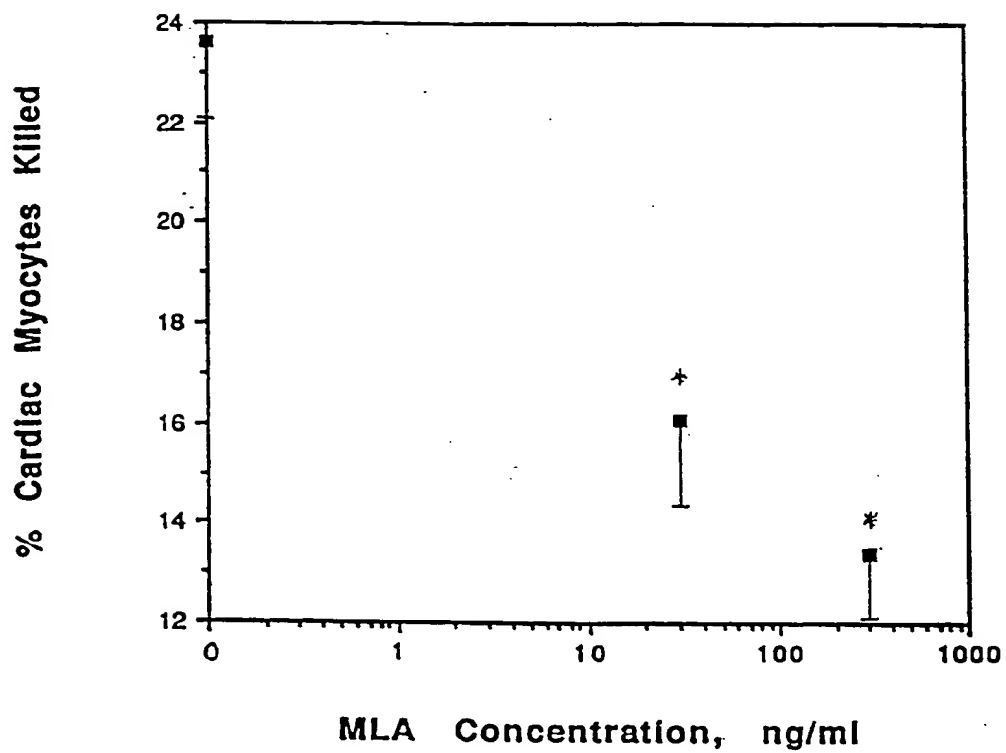


Figure 1

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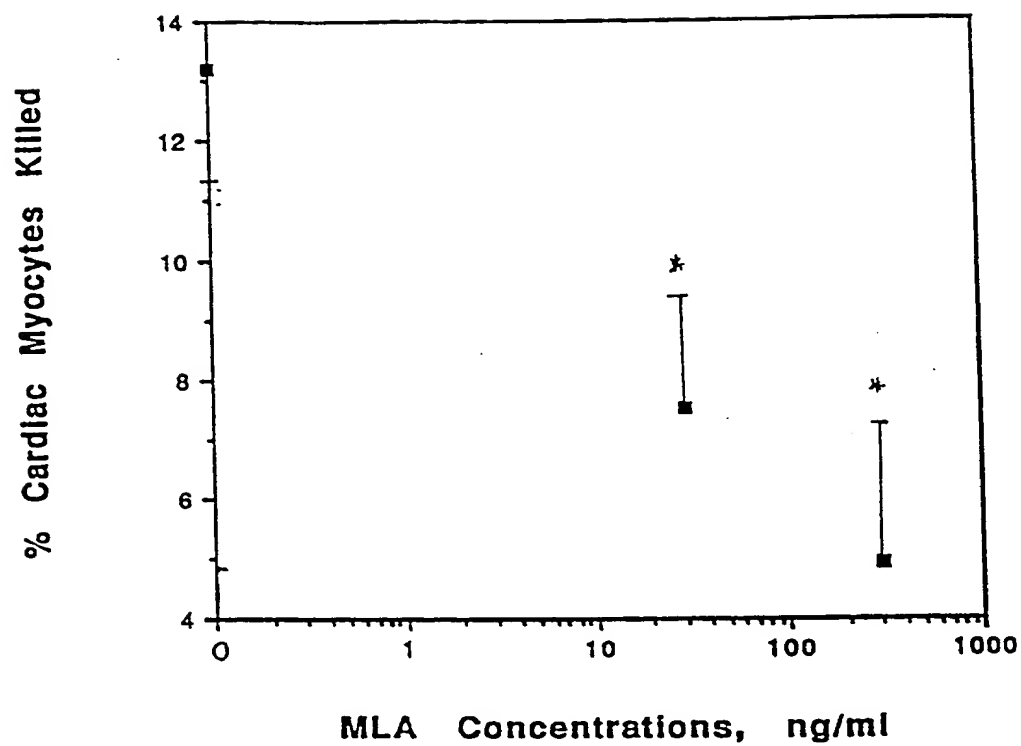


Figure 2

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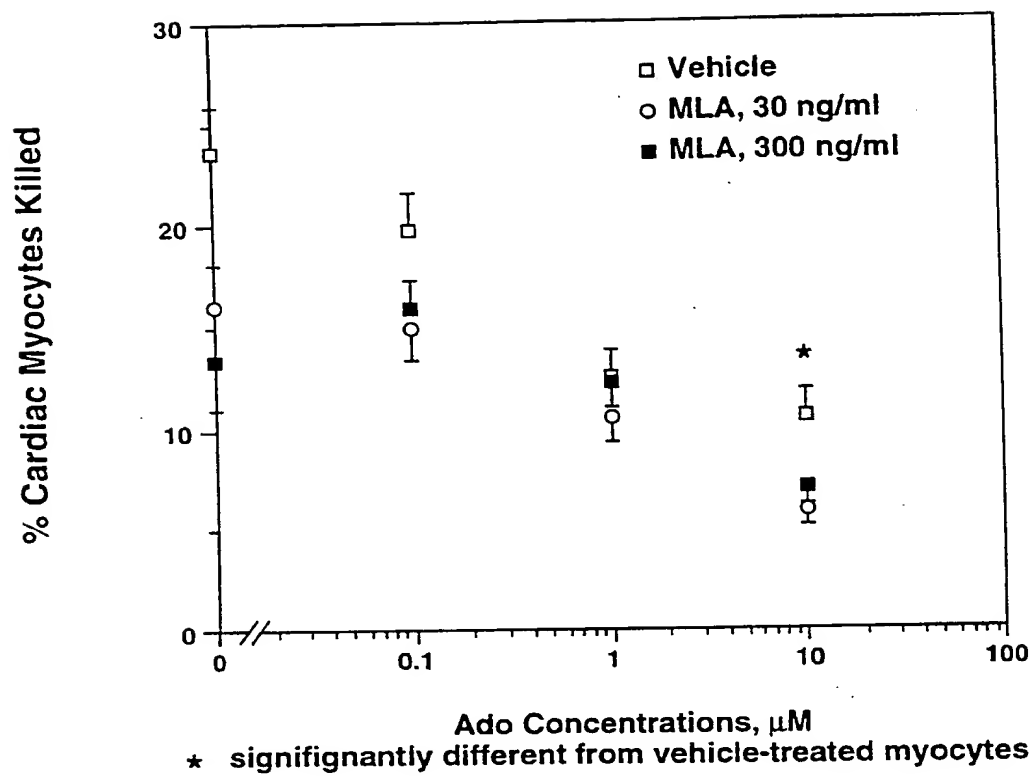


Fig. 3A

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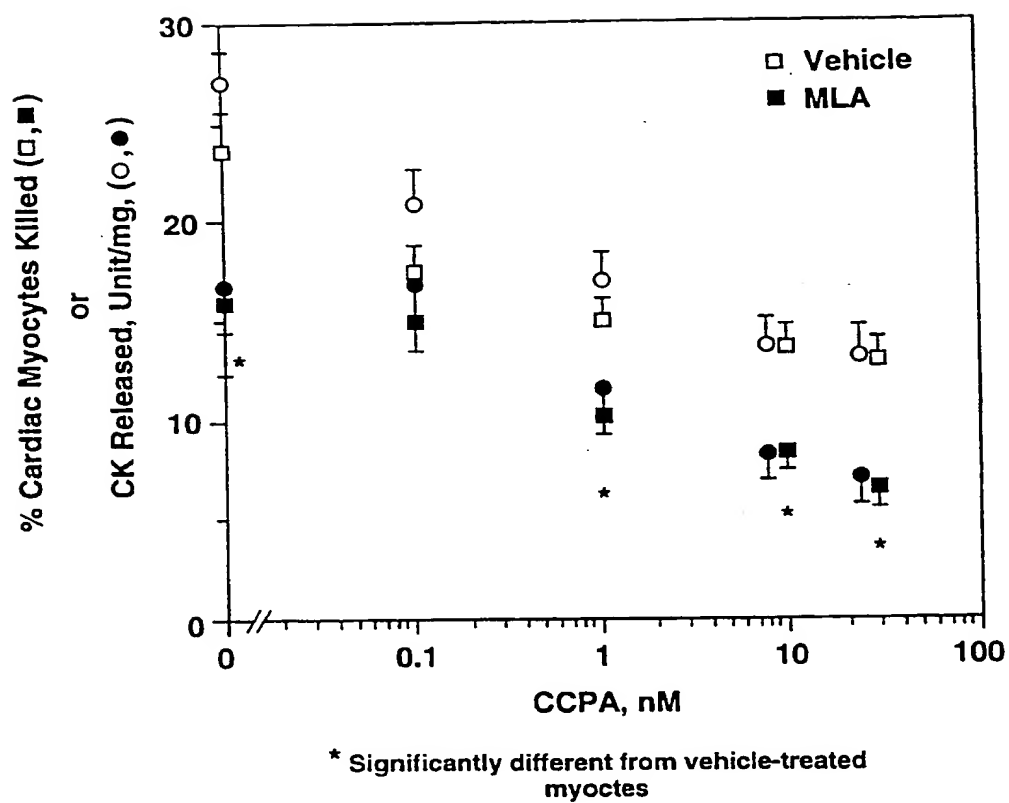
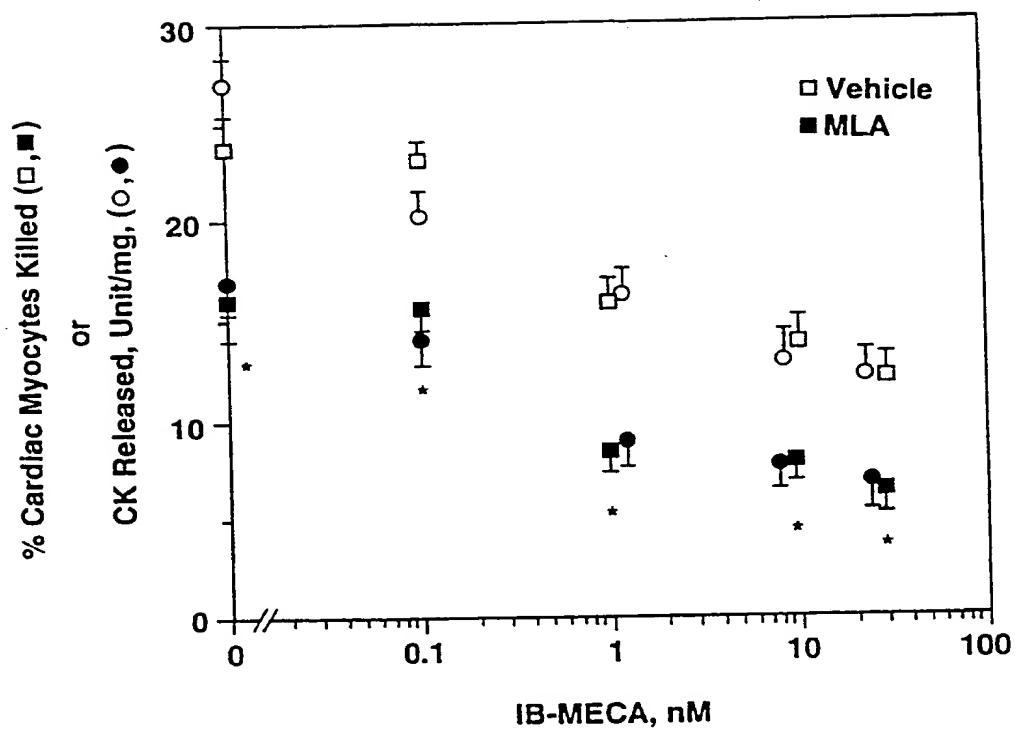


Fig. 3B

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* Significantly different from vehicle-treated myocytes

Fig. 3C

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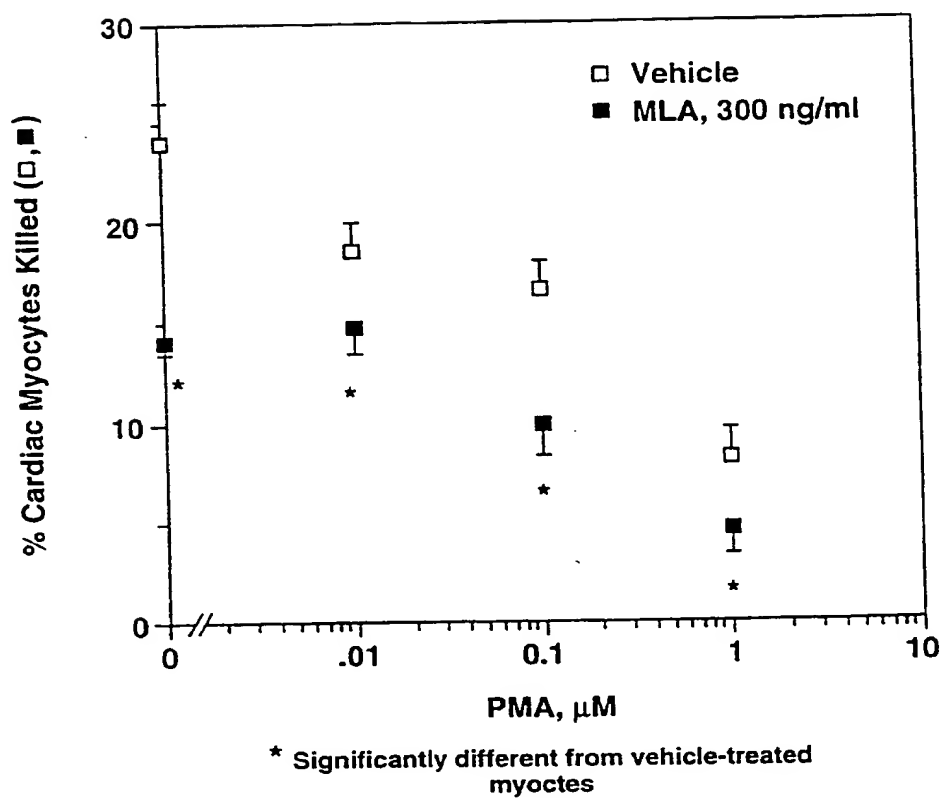


Figure 4

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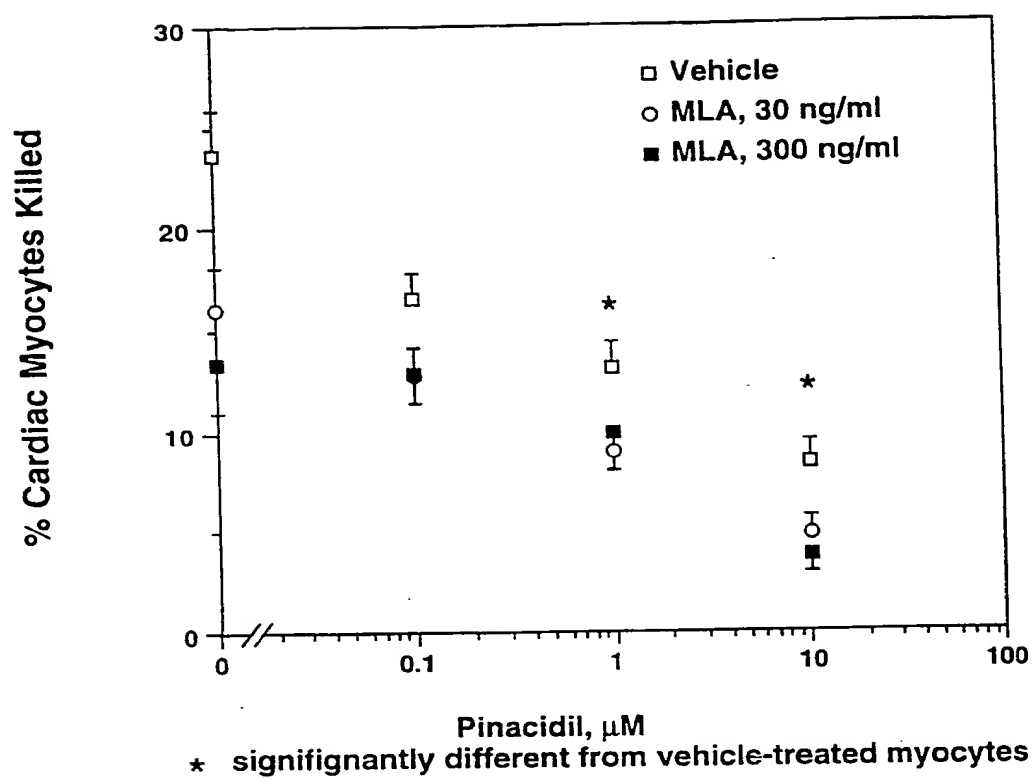
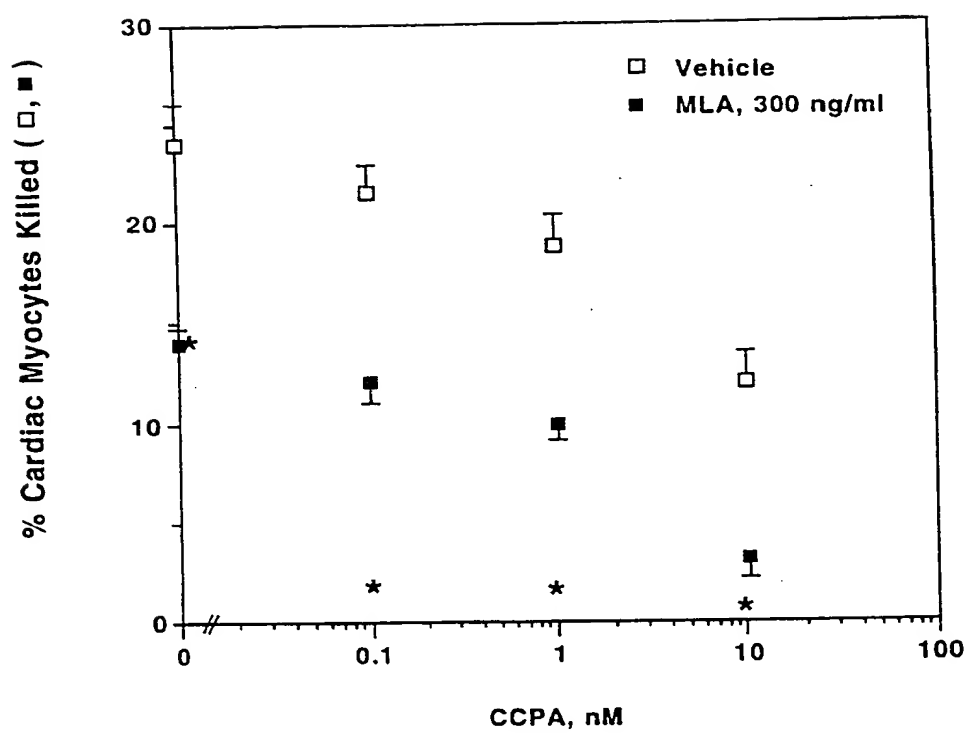


Figure 5

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* significantly different from vehicle-treated myocytes

Figure 6

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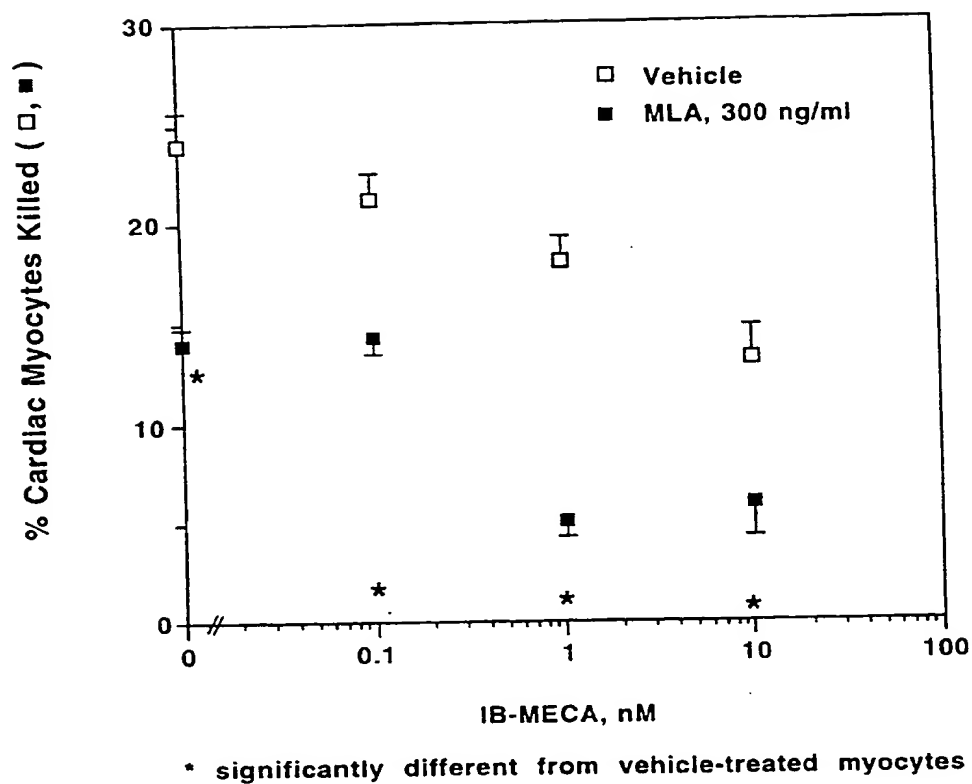


Figure 7

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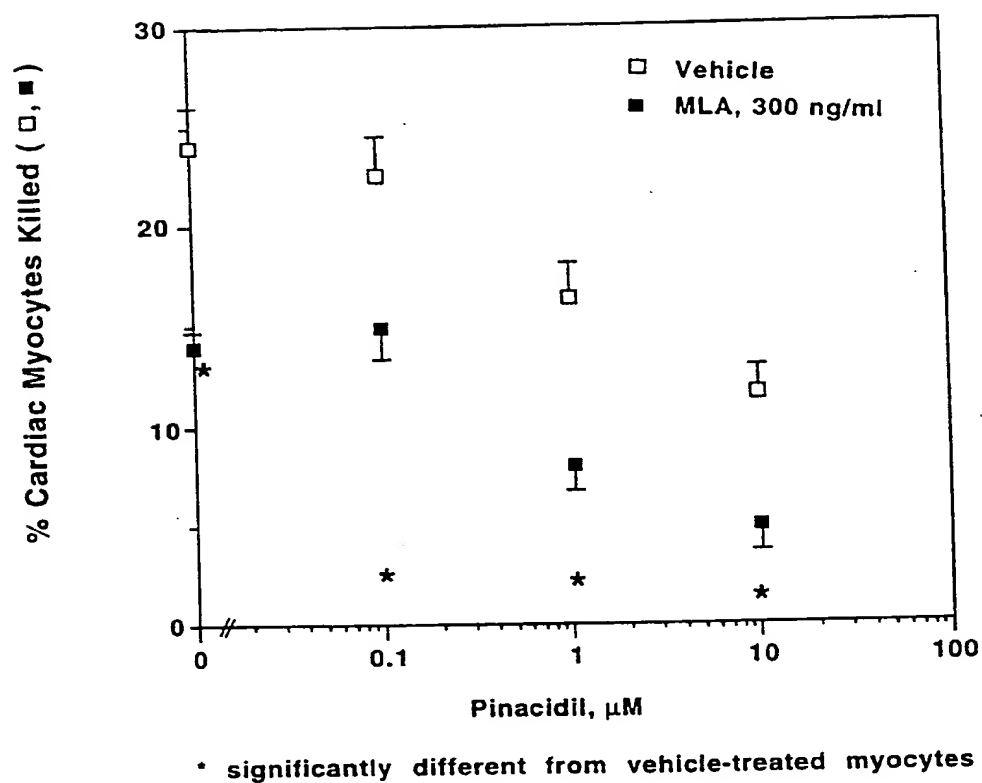


Figure 8

INTERNATIONAL SEARCH REPORT

International application No.
PCT/US98/22515

A. CLASSIFICATION OF SUBJECT MATTER

IPC(6) : A61K 31/715, 31/44
US CL : 514/54, 353, 355, 356, 921
According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)
U.S. : 514/54, 353, 355, 356, 921

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched
NONE

Electronic data base consulted during the international search (name of data base and, where practicable, search terms used)
APS, CAS-ONLINE

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
Y	US 4,870,090 A (NODA et al.) 26 September 1989, see the entire document.	1, 3, 5-10, 12, 14-24, 26, 28-30
Y	US 5,286,718 A (ELLIOTT) 15 February 1994, see the entire document.	1, 2, 4, 6-11, 13, 15-21, 23, 25-27, 29 and 30
Y	US 5,354,764 A (GROVER et al.) 11 October 1994, see the entire document.	1, 2, 4, 6-11, 13, 15-21, 23, 25-27, 29 and 30

☐ Further documents are listed in the continuation of Box C. ☐ See patent family annex.

* Special categories of cited documents:	*T* later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principles or theory underlying the invention
A document defining the general state of the art which is not considered to be of particular relevance	*X* - document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone
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L document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)	*G* document member of the same patent family
O document referring to an oral disclosure, use, exhibition or other means	
P document published prior to the international filing date but later than the priority date claimed	

Date of the actual completion of the international search
12 FEBRUARY 1999

Date of mailing of the international search report
19 FEB 1999

Name and mailing address of the ISA/US
Commissioner of Patents and Trademarks
Box PCT
Washington, D.C. 20231
Facsimile No. (703) 305-3230

Authorized officer
KEVIN E. WEDDINGTON
Telephone No. (703) 308-1235